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Harnessing mass differential confinement effects in magnetized rotating plasmas to address new separation needs

R. Gueroult

Laplace, Université de Toulouse, CNRS, INPT, UPS, 31062 Toulouse, France

E-mail: renaud.gueroult@laplace.univ-tlse.fr

J.-M. Rax

Université de Paris XI - Ecole Polytechnique, LOA-ENSTA-CNRS, 91128 Palaiseau, France

S. J. Zweben, N. J. Fisch

Princeton Plasma Physics Laboratory, Princeton University, Princeton, NJ 08543 USA

Abstract. The ability to separate large volumes of mixed species based on atomic mass appears desirable for a variety of emerging applications with high societal impact. One possibility to meet this objective consists in leveraging mass differential effects in rotating plasmas. Beyond conventional centrifugation, rotating plasmas offer in principle additional ways to separate elements based on mass. Single ion orbits show that ion radial mass separation in a uniform magnetized plasma column can be achieved by applying a tailored electric potential profile across the column, or by driving a rotating magnetic field within the column. Furthermore, magnetic pressure and centrifugal effects can be combined in a non-uniform geometry to separate ions based on mass along the field lines. Practical application of these separation schemes hinges on the ability to produce the desirable electric and magnetic field configuration within the plasma column.

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1. Introduction

New and innovative separation techniques could prove extremely valuable in a variety of applications. For instance, it was recently shown that substituting membrane-based separation for distillation in separation processes could lead to a 7% decrease of the total U. S. energy consumption [1]. The same study projects that implementing energy efficient separation techniques in the U. S. petroleum, chemical and paper manufacturing sectors alone could save 100 million tonnes of carbon dioxide emissions and \$4 billion in energy costs annually. Besides the economical incentive, the development of innovative separation techniques is also motivated by their anticipated enabling role in many applications [2, 3].

Physical separation techniques rely on differences in physical properties. Common examples are distillation, centrifugation and filtration, for which differences in respectively boiling point, mass and size are used. In essence, physical separation techniques harness differential transport and equilibrium properties in a species mixture. A sub-group of physical separation techniques is plasma separation techniques, where the feed to be separated is first turned into a plasma [4]. By ionizing the input feed, separation is carried out at the elemental level. As a result, the whole range of plasma transport phenomenon can in principle be leveraged to produce the desired separation properties.

Physical separation at the elemental level can be traced back to Dempster's mass spectrometer [5] and the calutron device [6], in which magnetic deflection was used to separate ions based on mass. In these devices, throughput is limited both by space charge effects [7, 8] and instabilities [9]. Since plasmas offer a natural neutralization mechanism, plasmas began being considered for separation applications. In particular, the realization that diffusion in a multi-ion species plasma subjected to centrifugal or gravitational forces exhibits asymmetrical effects [10–13] led to the development of plasma centrifuges [14–16] which were then used for isotope separation [17–20]. In these devices, collisional drag between species leads to an inward drift of the light species and an outward drift of the heavy species [10]. Separation arises from the mass dependent, and therefore species dependent, radial equilibrium density profile controlled by rotation [13, 21]. Although plasma centrifuges are conceptually similar to gas and liquid centrifuges, rotation in these

devices results from electromagnetic forces and not from frictional entrainment by moving parts. This difference allows for much larger rotation speeds, which in principle translates to higher separation power per centrifuge [22].

Besides the differential collisional drag exploited in plasma centrifuges, various other differential mechanisms in plasmas were proposed and studied to separate isotopes [23]. For example, differences in excitation energy were used in atomic vapor and molecular laser separation [24], while ionwave interactions, such as ion cyclotron resonance (ICR) [25,26], ponderomotive force [27,28] and hybrid resonance [29] were suggested for the development of electromagnetic separators. It is worth noting that both the ICR process through the TRW program [30] and the laser separation process through the AVLIS [31] and MLIS [32] programs were demonstrated in laboratory at large scale. However, a large part of the research work conducted on this topic has been either classified or proprietary in the past (e. g. the calutron and the TRW work), which has made it hard to obtain a comprehensive summary of past work.

Isotope separation stands out from other separation needs owing to the small mass difference between the elements to be separated. A legitimate question is therefore to ask whether new plasma mechanisms can be put forward to efficiently separate elements by relaxing the constraint on the mass difference. In this paper, we offer some perspectives on this question by introducing theoretical separation concepts and estimates. However, none of these ideas has yet been implemented in a practical industrial-scale production device. First, in Sec. 2, a variety of applications with high societal impact and for which high-throughput plasma mass separation could prove valuable is highlighted. In light of this observation, in Sec. 3, mass differential confinement properties for a particular class of configurations, namely rotating plasmas, are reviewed. The mass separation potential of rotating plasmas is first considered in a uniform axial magnetic field, and then extended to non-uniform fields. In Sec. 4, the main findings are summarized.

2. Rationale for developing high-throughput plasma mass separation techniques

2.1. Need for new separation technologies

Owing to the role they play in many industries, e. g. chemical, petroleum refining and materials processing, and the opportunities they present for waste reduction and energy efficiency, the development of separation technologies is of great importance (see, e. g., Refs. [2, 3]).

For example, consider nuclear waste cleanup in the United States [33], which is projected to cost more than 280 billion dollars over the next 40 years [34]. Cleanup effort requires separating high-activity waste from low-activity waste before vitrification and eventual storage in a geological repository. Due to the heterogeneous and dynamic properties of some of these wastes, conventional chemical separation techniques are particularly challenged. In the meantime, the cleanup cost depends strongly on the efficiency of waste partitioning and minimization [35]. Therefore, "the development of robust and well-understood technologies to enable safe, selective, efficient, and cost-effective cleanup of wastes" has recently been called for [36].

Another example is nuclear spent fuel reprocessing [37, 38] as envisioned in advanced nuclear fuel cycles [39,40]. Spent fuel reprocessing and closed fuel cycles, through partitioning and transmuting long lived actinides into shorter lived elements, can decrease the lifetime and associated biological hazards of nuclear spent fuel to a few hundred years [41, 42]. However, transmutation requires separating actinides from lanthanides found in spent fuel because of the larger neutron capture cross section of lanthanides [43]. Owing to the similar chemical properties of these elements [44, 45], separating lanthanides (4-f block elements) from actinides (5-f block elements) requires multiple complex chemical stages, which comes at the expense of cost and reliability [46]. Therefore, a single stage process is desirable.

Finally, yet another example is rare earth recycling. Expansion of rare earth recycling could mitigate the risks associated with the high-volatility of rare earth market [47], as well as limit the environmental impact associated with rare earth mining [48]. However, existing hydro-metallurgical recycling pathways are very similar to the multiple stage processes used for extraction from primary ores [49], and hence also often suffer from a significant environmental footprint. Here again, a clean single stage process is desirable.

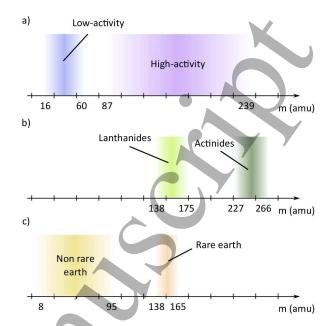


Figure 1. Composition of the input feed as a function of atomic mass for various separation needs: a) separation of high activity waste from low activity waste in nuclear waste cleanup, from [50], b) actinides/lanthanides separation in nuclear spent fuel reprocessing, from [51] and c) rare earth separation in rare earth recycling of NdFeB magnets, from [52].

2.2. Interest for plasma mass separation

Looking at the elemental composition of the feed and desired product streams of these three separation needs reveals an interesting characteristic. As shown in Fig. 1, the components to be separated break down into a light and a heavy component for each application. For nuclear waste cleanup (Fig. 1a), high-activity elements are much heavier than lowactivity elements [50]. For spent fuel reprocessing (Fig. 1b), lanthanides are lighter than actinides [51]. Finally, for rare earth recovery from NdFeB magnets (Fig. 1c), rare earth elements are heavier than all other constituents [52]. In all three cases, the light and heavy components are separated by a gap of 30-50 atomic mass units. In light of this feature, it appears that mass separation at the elemental level could prove valuable for these three applications.

Besides the ability to separate elements adequately, another constrain lies in the ability to process the amount of material required by a given application. For example, take spent fuel reprocessing. The largest chemical-based nuclear reprocessing plants handle some 10^6 kg of spent fuel per year, of which actinides and lanthanides account for about 10^3 and 10^4 kg, respectively [51,53]. Separation of actinides from lanthanides will thus require a throughput of about 10^4 kg per year.

In plasma mass separation devices designed

for isotope separation, a trade-off generally exists between throughput and enrichment factor as a result of collisions. In the TRW experiment [30], the largest experimental throughput produced was about 10 kg/year, and the best projections were of the order of 100 kg/year [54, p. 123]. enrichment factor would benefit from larger mass difference allowing in turn for larger throughput, this is unlikely to suffice to meet the requirement of, say, spent fuel reprocessing. Concepts combining ICR selective heating with particle drift in curved magnetic field could possibly achieve 10³ kg/year [55], but they are still limited by collisions. There is therefore an incentive to look for new plasma mass separation schemes which would allow high-throughput processing.

2.3. Cost of plasma separation

Although mass separation at the elemental level holds promise for the applications discussed at the beginning of this section, it remains to be shown that these processes are practically feasible and that the cost of plasma separation is not prohibitive.

After a possible pre-treatment operation, plasma separation first requires turning the input feed into a plasma. In practice, this step breaks down into two sub-steps. The material is first heated to become a gas in the evaporation step. Upon further heating, the gas is turned into a plasma in the ionization step. Once the input feed is ionized, the plasma has to be maintained while separation processes occur. An estimate of the lower limit of the separation cost can be derived based on the cost of each of these processes.

The first process consists in turning the input feed into a plasma. Strictly speaking, the energy cost for turning solid material into a gas is the sum of the latent heat of both fusion and vaporization, plus the enthalpy change corresponding to heating the material from room temperature to melting temperature, and then from melting temperature to boiling temperature. However, the latent heat of vaporization \mathcal{L}_V typically dominates the other contributions, and the energy cost of turning material into a gas is in first approximation \mathcal{L}_V . For metals, $\mathcal{L}_V \sim 1-10$ MJ/kg. This however does not include any losses. Using laser ablation as a baseline, the real energy cost for evaporation depends on the laser absorptivity χ , which is typically 0.1 - 0.4 for common metals [56]. Accounting finally for the laser electric efficiency η_l , the energy cost for evaporation is $\eta_l^{-1}\chi^{-1}\mathcal{L}_V$. For an assumed $\eta_l=0.1$, this is at most 1 GJ/kg.

An estimate for the energy cost of ionization can be obtained by assuming a fully ionized plasma and a given chemical composition. Ionization energy for atoms varies between 3.8 eV for francium and 24.6 eV for helium. One kilogram of material of average atomic mass $m_i \sim 100$ amu is made of about 6 10^{24} atoms. The energy cost for fully ionizing 1 kg of such material is hence roughly 3-25 MJ/kg. Here again, energy losses such as excitation and radiation losses need to be accounted for. For helicon plasmas, the efficiency of plasma formation η_p has been shown to be about 0.4 in pure argon [57]. Since complex plasmas will add extra energy dissipation channels, the plasma efficiency is expected to be lower in this case. For a very degraded $\eta_p = 0.02$, the cost of plasma formation and and maintenance is about 0.15-1.25 GJ/kg.

Summing these two contributions, an upper bound energy cost for plasma separation is 2 GJ/kg. For an electricity cost of \$0.12 per kW.h, this is about \$65 per kg. It is worth pointing out here that this figure could be significantly lower if one could produce separation in a partially ionized plasma since the ionization cost scales with the number of ionized atoms. Besides processing costs considered thus far, capital, operation and maintenance costs will have to be accounted for. However, owing to a comparatively small footprint, capital costs of plasma techniques are expected to be lower than those of chemical techniques. This scaling was for example observed when comparing high-temperature processing and aqueous processing [58].

A processing cost of \$65 per kg already suggests that plasma techniques, at least in their current form, are unlikely to be attractive for applications for which proven techniques are readily available. However, preliminary cost comparative suggests that plasma techniques might be competitive with the proposed chemical solutions for nuclear waste cleanup thanks to improved waste minimization [50]. Similar considerations suggest that plasma processing of NdFeB magnets for rare earth recovery could be economically attractive [52]. Finally, the continuous plasma processing of spent fuel unloaded from a nuclear reactor has been estimated to only require 0.06% of the output power of this reactor [59].

Further to this point, a complete cost comparison should include the environmental, social and economical costs and benefits of each separation techniques. Although these effects are difficult to quantify, this is where plasma techniques could prove particularly advantageous. It may be that the very limited environmental cost of plasma techniques will offset a possible disadvantage when considering processing costs alone.

3. Mass differential confinement effects in magnetized rotating plasmas

In the search for mass differential effects, rotating configurations hold particular promise thanks to the centrifugal effects associated with rotation. In the

remaining of this paper, we therefore analyze in a systematic manner mass differential confinement properties in magnetized rotating plasmas.

3.1. $E \times B$ rotating plasmas in purely axial magnetic field

Two relatively simple configurations can be readily identified to produce rotation as a result of the $\mathbf{E} \times \mathbf{B}$ drift in a plasma column: an axial magnetic combined with a radial electric field, or, alternatively, a radial magnetic field and an axial electric field. The former of these two configurations, with a uniform axial magnetic field $\mathbf{B} = B_0 \mathbf{e_z}$ and a radial electric field $\mathbf{E} = E_r \mathbf{e_r} = -\nabla(\phi)$ as depicted in Fig. 2, displays the interesting property that the cross-product of the centrifugal force with the magnetic field is non zero. As a result, centrifugal forces cause an additional azimuthal drift.

Considering the plasma column in Fig. 2, and neglecting first collisions, the radial force balance on a particle of charge q and mass m writes

$$-\omega^2 = \frac{q}{mr} E_r + \operatorname{sgn}(q) \Omega \omega, \tag{1}$$

with ω the azimuthal angular frequency of the gyrocenter, $\operatorname{sgn}(x) = x/|x|$ the sign function and $\Omega = |q|B_0/m$ the cyclotron frequency. The equilibrium solution is described by the slow and fast Brillouin modes [60]

$$\omega_B^{\pm} = -\operatorname{sgn}(q) \frac{\Omega}{2} \left[1 \pm \sqrt{1 - \frac{4mE_r}{qB_0^2 r}} \right]. \tag{2}$$

Out of these two modes, only the slow mode ω_B^- depicted in Fig. 3 arises spontaneously. Introducing the azimuthal drift velocity in the limit of zero inertia $\Omega_E = -E_r/(rB_0)$, Eq. (2) rewrites

$$\omega_B^- = -\operatorname{sgn}(q)\frac{\Omega}{2}\left[1 - \sqrt{1 + 4\operatorname{sgn}(q)\frac{\Omega_E}{\Omega}}\right].$$
 (3)

Taylor expanding Eq. (3) for $|\Omega_E|/\Omega \ll 1$, one gets

$$\omega_B^- = \Omega_E \left[1 - \operatorname{sgn}(q) \frac{\Omega_E}{\Omega} + \mathcal{O}\left(\left[\frac{\Omega_E}{\Omega} \right]^2 \right) \right]$$
 (4)

In the limit $|\Omega_E|/\Omega \to 0$, one recovers $\omega = \Omega_E$. In this limit, there is no difference in azimuthal $\mathbf{E} \times \mathbf{B}$ drift velocity between charged species.

For $E_r \leq 0$, $\Omega_E \geq 0$, and plasma rotation is in the counter-clockwise direction. Here, clockwise is defined for the case where the magnetic field is pointing out of the page, that is to say the negative $\hat{\theta}$ direction. Eq. (3) shows that centrifugal effects speed up rotation for negatively charged particles, and slow up particles for positively charged particles, as seen in Fig. 3. For two positive ions of different mass, the angular velocity of the light ion is larger than the angular velocity of the heavy ion. On the other hand, rotation is in the clockwise direction for $E_r \geq 0$, and $\Omega_E \leq 0$. For this polarity, positively charged particles rotate faster, while negatively charged particles rotate slower. For two positive ions of different mass, the norm of the angular velocity of the light ion is smaller than the norm of the angular velocity of the heavy ion. Note that the difference in azimuthal velocity between two different ion species leads to a positive (resp. negative) azimuthal drag force on heavy (resp. light) ions no matter the polarity of the radial electric field. In both cases, this drag force causes light ions to drift radially inward and heavy ions to drift radially outward. This is the physical mechanism behind plasma centrifugation [11].

Now, looking at Eq. (3), one notices that there is no solution if

$$\operatorname{sgn}(q)\frac{\Omega_E}{\Omega} \ge -1/4. \tag{5}$$

This limit, known as the Brillouin limit, means that ions are radially unconfined for fast enough rotation in the clockwise direction, and electrons are unconfined for fast enough rotation in the counter-clockwise direction. Since $\Omega_e \gg \Omega_i$, the latter is however unlikely.

To illustrate these confinement properties, it is interesting (as it will become clear later) to consider particle equilibrium in the frame rotating with the angular velocity $\boldsymbol{\varpi} = -\mathrm{sgn}(q)\Omega/2~\mathbf{e_z}$. Let us denote variables in this rotating frame with a $\tilde{}$. Since $\partial \varpi/\partial t = 0$, the fields transformation reads

$$\tilde{\mathbf{E}} = \mathbf{E} + (\boldsymbol{\varpi} \times \tilde{r}) \times \mathbf{B} \tag{6a}$$

$$\tilde{\mathbf{B}} = \mathbf{B}.\tag{6b}$$

Assuming the fields in the rotating frame do not depend on time, one can rewrite the Newton-Lorentz equation as (see for example, Ref [61, p. 328])

$$m\frac{\partial \tilde{\mathbf{v}}}{\partial t} = q(\mathbf{E}^* + \tilde{\mathbf{v}} \times \mathbf{B}^*) \tag{7}$$

with

$$\mathbf{E}^{\star} = \tilde{\mathbf{E}} + \mathbf{\nabla} \left(\frac{m \varpi^2 \tilde{r}^2}{2q} \right) \tag{8a}$$

$$\mathbf{B}^{\star} = \tilde{\mathbf{B}} + \frac{2m}{q}\boldsymbol{\varpi} = 0. \tag{8b}$$

Eq. (8b) shows that in the chosen frame rotating with the angular frequency $\boldsymbol{\varpi} = -\mathrm{sgn}(q)\Omega/2 \,\mathbf{e_z}$, the magnetic field cancels. In this frame, the particle dynamics is only controlled by the electric field \mathbf{E}^{\star} . The second term on the right hand side in Eq. (8a) is the contribution of the centrifugal force. Introducing the effective potential

$$\phi^{\star}(\tilde{r}) = \phi(r) + [2\operatorname{sgn}(q) - 1] \frac{qB_0^2}{8m} \tilde{r}^2, \tag{9}$$

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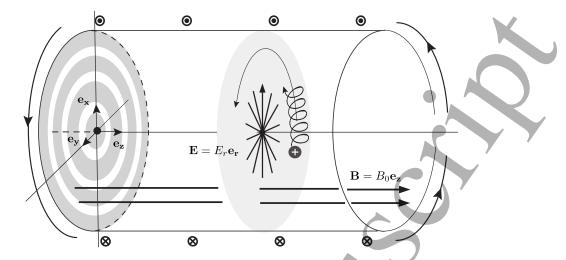


Figure 2. Linear configuration: uniform axial magnetic field and radial electric field.

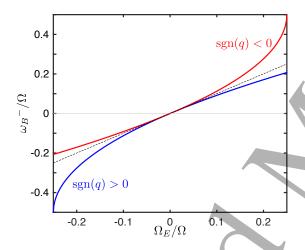


Figure 3. Slow Brillouin mode for positively (blue) and negatively (red) charged particles, with $\Omega = |q|B_0/m$ the gyrofrequency and $\Omega_E = -E_r/B_0$ the $\mathbf{E} \times \mathbf{B}$ drift angular frequency. Rotation is counter-clockwise ($\omega_B^- > 0$) for $E_r < 0$, and reciprocally. The black dotted curve represents the zero inertia solution.

Eq. (8a) writes $\mathbf{E}^* = -\nabla \phi^*$. Since the Coriolis force is proportional to ϖ , it depends on the sign of the rotation and therefore here on the sign of the particle charge. In contrast, the centrifugal force is proportional to ϖ^2 , and is therefore positive irrespective of the sign of the particle charge. For positively charged particles, Coriolis and centrifugal forces are in opposite direction, and one gets

$$\phi^{\star}(\tilde{r}) = \phi(r) + \frac{qB_0^2}{8m}\tilde{r}^2. \tag{10}$$

If the potential applied in the laboratory frame is constant $(\partial \phi/\partial r = 0)$, the effective potential ϕ^* in Eq. (10) is convex, and ions are confined. Eq. (9) shows electrons are also confined in this case. Now assume a

parabolic potential profile $\phi(r) = \alpha r^2$ is applied in the laboratory frame. This corresponds to a solid body rotating plasma column since $E_r \propto r$ so $\partial \Omega_e/\partial r = 0$, and thus, using Eq. (3), $\partial \omega_B^-/\partial r = 0$. For $\alpha \geq -qB_0^2/(8m)$, an ion of mass m and charge q is still confined. On the other hand, for $\alpha \leq -qB_0^2/(8m)$, Eq. (10) tells us that ϕ^* is concave. An ion of mass m and charge q is therefore radially unconfined. The change in concavity of the effective potential profile $\phi^*(r)$ is illustrated in Fig. 4. The threshold value $\alpha_c = -qB_0^2/(8m)$ for ion confinement can be rewritten $E_r/(rB_0) = \Omega/4$, which is the Brillouin limit given in Eq. (5). Now suppose a multi-ion species plasma with $\alpha = -qB_0^2/(8m_{\phi})$, so that

$$\phi^{\star}(\tilde{r}) = \frac{qB_0^2}{8mm_{\diamond}}(m_{\diamond} - m)\tilde{r}^2. \tag{11}$$

The effective potential ϕ^* indicates that a singly charged ion with mass $m \geq m_{\diamond}$ will be radially unconfined, while a singly charged ion with mass $m \leq m_{\diamond}$ will be radially confined. This charge to mass ratio threshold for confinement is the basis for the DC band gap ion mass filter [62] used in the Archimedes plasma mass filter [63]. In this device, ions are separated into two components: light ions $m/m_{\diamond} < 1$ are collected axially along the magnetic field lines while heavy ions $m/m_{\diamond} > 1$ are collected radially.

Practically, this filtering mechanism has a few limitations. First, since the confinement criteria depends on the charge to mass ratio and not on the mass alone: a doubly charged ion of mass 2m can not be differentiated from a singly charged ion of mass m. This means that heavy doubly charged ions will be collected with light singly charged ions. Second, the filtering mechanism relies on low collisionality, which sets a limit on plasma density and hence throughput for a given magnetic field intensity. Indeed, the

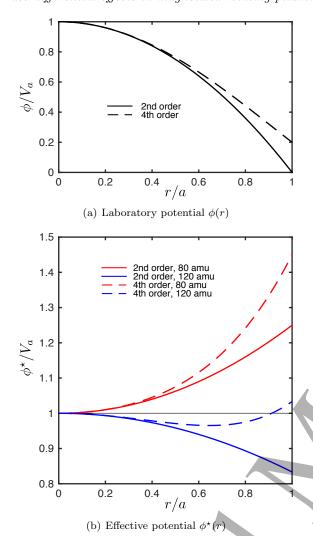


Figure 4. Applied laboratory potential radial profile $\phi(r)$ [(a)] and effective potential radial profile $\phi^*(r)$ for two different ion mass [(b)]. Solid line curves are obtained for $\phi(r) = V_a(1-r^2/a^2)$, while dotted line curves are obtained for $\phi(r) = \beta r^4 + V_a(1-r^2/a^2)$, $\beta \in \mathbb{R}$. V_a is the potential difference across the plasma column for the parabolic case, a is the plasma column radius. The addition of a fourth order term to the parabolic profile leads to the formation of a potential well off-axis for heavy ions (dotted blue curve).

radial ion transport induced by collisions with neutrals brings light ions $(m/m_{\diamond} < 1)$ to the heavy ions $(m/m_{\diamond} > 1)$ stream. Strictly speaking, ion-neutral collisions slow down the slow mode and suppress the requirement for ion radial confinement $\Omega_E/\Omega \ge -1/4$ [64]. In other words, the Brillouin limit breaks down. Separation then hinges on the differential radial transport properties of light and heavy ions. Finally, and maybe most importantly, another limitation is that heavy ions are typically collected over a large region of the plasma chamber. This is because heavy ions are extracted perpendicularly to the field lines

with little control over their deposition and extraction. This is particularly an issue when heavy particles are made of hazardous materials, for example for nuclear waste cleanup.

To remediate to this last issue, one can use a higher order polynomial profile for the laboratory potential ϕ . For example, a fourth order polynomial can be used to create a dip in effective potential offaxis, while maintaining global radial confinement [65]. This scheme is illustrated in Fig. 4. lighter than the mass threshold (red dotted curve in Fig. 4(a)), the effective potential ϕ^* is monotonically increasing with r, so that light ions are collected axially along the field lines in the central region. On the other hand, heavy ions see a minimum in effective potential off-axis (blue dotted curve in Fig. 4(b)), but are still radially confined as opposed to the parabolic potential case. Heavy ions are therefore collected axially along the field lines but in an annular region at larger radius. In contrast with the DC band gap ion mass filter [62] for which collisions are detrimental to separation performances, collisions are in this configuration required since they allow radial diffusion of heavy ions towards the off-axis potential well. Without collisions, heavy ions would extend radially from the center to a point r_h past the minimum of ϕ^* and which depends on the ion temperature. Radial separation of heavy from light ions can be optimized through the radial profile of the laboratory potential $\phi(r)$ and the device geometry [65]. However, the advantage of extracting both light and heavy species along field lines comes at the expense of producing and controlling a more complex potential radial profile in the plasma. Also, the use of higherorder potential profile means that plasma rotation is now sheared $(\partial \omega/\partial r \neq 0)$, which is known to lead, under certain conditions, to the onset of Kelvin-Helmholtz instabilities [66, 67].

The difference in suitable collisionality regime also provides insights into the potential of each scheme for high-throughput separation. Collisionless operation in the DC band gap ion mass filter [62] will limit the plasma density and hence the achievable throughput for a given magnetic field. On the other hand, no such limit on plasma density is found in the higher-order scheme [65]. This result suggests that higher-order schemes might be advantageous for high-throughput separation.

To conclude this discussion of mass separation due to $\mathbf{E} \times \mathbf{B}$ rotation in a uniform magnetic field, it is worth noting that both radial-axial mass separation in Archimedes filter [63] and radial-radial mass separation in the double well mass filter [65] require fast plasma rotation. Quantitatively, separation occurs near the Brillouin limit for which $|\omega|/\Omega \sim 1/2$. For this rotation

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regime, significant differences in azimuthal velocities can exist between ions with different mass, which could trigger the onset of centrifugal instabilities [68–70]. In light of this observation, it seems advantageous if possible to produce mass separation at lower rotation velocity. This might be done by abandoning the uniform axial magnetic field topology considered up to this point.

3.2. $E \times B$ rotating plasmas in inclined magnetic fields

Substituting an inclined magnetic field $\mathbf{B} = B_r \mathbf{e_r} + B_z \mathbf{e_z} = B \mathbf{e_b}$ in place of the purely axial field considered in the previous section offers additional means of control. Assume a conical magnetic surface defined by $\mathbf{B} \cdot \mathbf{e_z} = \cos \alpha$, and write s the curvilinear coordinate along a given field line. Flux conservation requires $B(s) = B_0 r_0 / r(s)$, where B_0 and r_0 are the magnetic field intensity and the field line radius at s = 0, respectively. In this configuration, the magnetic mirror force

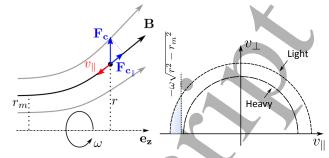
$$-\mu \frac{\partial B}{\partial s} \mathbf{e_b} = \mu B_0 \frac{r_0}{r(s)^2} \sin \alpha \mathbf{e_b}, \tag{12}$$

with $\mu=mv_{\perp}^2/(2B)$ the magnetic moment of the particle. Besides this mirror force, the force balance along the field line requires accounting for the contribution of centrifugal forces produced by plasma rotation $m\omega^2 r(s)\sin\alpha\mathbf{e_b}$, with ω the rotation velocity. Note that the iso-rotation theorem states that ω is constant on a given magnetic surface (see, e. g., Refs. [12,71]). Centrifugal and mirror forces add up, and a particle moving along the field line towards larger r accelerates, while a particle moving towards smaller r slows down.

Consider now the field topology depicted in Fig. 5(a). A particle at radius r with negative v_{\parallel} sees a centrifugal potential barrier $m\omega^2(r^2-r_m^2)/2$. Interestingly, this potential barrier is proportional to the particle mass. For a given parallel energy ϵ_{\parallel} and two particles of mass m_l and m_h with $m_l \leq m_h$, there therefore exists a rotation velocity ω for which the light particle can reach r_m , while the heavy particle can not. Assuming a two-ion species plasma in thermal equilibrium, this result can in principle be used to preferentially collect light ions on the left side in Fig. 5(a), as illustrated in Fig. 5(b). Strictly speaking, one should also consider the mirror force which tends to pull particles to larger r. However, from Eq.(12), the ratio of centrifugal to mirror forces along the field line is

$$\frac{\mathbf{F_c} \cdot \mathbf{e_b}}{-\mu \partial B / \partial s} = \frac{2r^3 \omega^2}{r_0 v_{\perp_0}^2},\tag{13}$$

and mirror effects should be negligible for large enough r



(a) Centrifugal end plug created by an inclined magnetic field

(b) Collection diagram at r_m for heavy and light ions in thermal equilibrium

Figure 5. Magnetic field topology [(a)] and mass separation capabilities [(b)] of a centrifugal end plug. The blue shaded region in Fig. (b) represents the part of ions starting at a radius r which are collected at the minimum radius r_m along the same field line. Mirror forces are here neglected.

Centrifugal and mirror effects can also be used in a way that they oppose each other, for example by creating a magnetic mirror at a larger radius as illustrated in Fig. 6(a). In this geometry, energy conservation yields the confinement criteria [12,72]

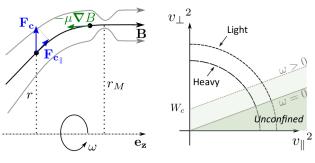
$$v_{\parallel}^{2} \le (r\Omega)^{2} \left(1 - \frac{r_{M}^{2}}{r^{2}}\right) + v_{\perp}^{2} \left(\frac{B_{M}}{B} - 1\right),$$
 (14)

where r_M and B_M are the field line radius and field intensity at the mirror, respectively. In the configuration depicted in Fig. 6(a), $r_M/r > 1$, so that the first term on the right hand side in Eq. (14) is negative, while the second term on the right hand side is positive. As a result, a particle with $v_{\parallel} = 0$ is only confined if $v_{\perp}^{-2} \ge W_c$, with

$$W_c = \left[\left(\frac{r_M}{r} \right)^2 - 1 \right] \left(\frac{B_M}{B} - 1 \right)^{-1} r^2 \omega^2. \tag{15}$$

This is in contrast with conventional rotating mirror machines $(r_M/r \ll 1)$ in which centrifugal forces combine with mirror forces to enhance ion confinement [73,74]. The threshold in perpendicular velocity described by Eq. (15) creates mass differential confinement properties as shown in Fig. 6(b). Considering again a two-ion species plasma in thermal equilibrium, the loss cone modified by rotation will cover a larger fraction of the distribution of heavy ions compared to the distribution of light ions. The rotation velocity ω provides control over the fraction of heavy ions lost through the mirror.

These two effects, namely preferential collection of light ions at smaller radius (Fig. 5) and preferential collection of heavy ions through a magnetic mirror at large radius (Fig. 6), are the basis of the Magnetic Centrifugal Mass Filter (MCMF) [75]. In this device, collisionality has to be large enough for ion-ion pitch angle scattering to scatter ions into the small radius side



(a) Mirror end plug at large (b) Particle confinement for heavy radius and light ions in thermal equilibrium

Figure 6. Magnetic field topology [(a)] and mass separation capabilities [(b)] of a mirror end plug. W_c is defined in Eq. (15). The shaded area in Fig. (b) represents the mirror loss cone, which grows with the rotation velocity ω .

loss cone, but low enough to limit perpendicular transport. The mass separation capabilities were confirmed through preliminary numerical simulations [51,76], and constrains imposed by collisionality on the operating window were recently highlighted [77]. However, and although collisionality sets an upper limit on practical density for a given ion temperature, projections appear compatible with high-throughput operation [51].

One critical question on which hinges the demonstration of the practicality of $\mathbf{E} \times \mathbf{B}$ rotating plasma configurations for mass separation is the ability to establish and control the required perpendicular electric field in the plasma. One possible solution is end electrodes biasing [78]. Although experiments in low density micro-wave and radio-frequency discharges provided encouraging results [79-81], further work is required to fully confirm the efficiency of this scheme at higher density [82]. Alternatively, wave-induced rotation has been suggested [83] as a way to suppress the need for end electrodes. This effect relies upon injecting waves with azimuthal phase velocities into the plasma to move charge across field lines [84]. In rotating plasma, this effect can create a radial potential that then drives the E x B flow [85]. While aspects of this effect have been demonstrated experimentally. the rotation effect has not yet been demonstrated in plasma devices suitable for separation. Another possibility to produce plasma rotation might lie in the use of rotating magnetic fields.

3.3. Plasma rotation in rotating magnetic fields

Alfvén's frozen in theorem predicts that a magnetized plasma column with an axial static magnetic field $B_0\mathbf{e_z}$ can, under certain conditions, be spun using a rotating magnetic field [86]. This configuration is depicted in

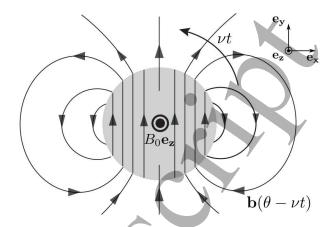


Figure 7. Linear configuration: static axial magnetic field $B_0 \mathbf{e_z}$ plus rotating magnetic field $\mathbf{b} = b_0 [\cos(\nu t) \mathbf{e_x} + \sin(\nu t) \mathbf{e_y}]$. The gray region represents the plasma column.

Fig. 7. However, single particle dynamics in rotating field configurations is far more convoluted than simple rotation [87-89]. It was also recently shown that ion confinement in rotating magnetic fields depends strongly on the particular external driving currents geometry used to produce this rotating magnetic field [90]. Here ion confinement is used in the sense of a stable ion orbit, that is to say that the ion radial position does not diverge. Examples of possible driving currents include a rotating axial current along $\mathbf{e}_{\mathbf{z}}$, a standing surface Alfvén wave along a cylindrical plasma column and two coils in quadrature and oriented perpendicularly (see Ref. [90]). The dependence of ion stability on the driving currents geometry can be understood by noting that the rotating magnetic field

$$\mathbf{b} = b_0 [\cos(\nu t) \mathbf{e_x} + \sin(\nu t) \mathbf{e_v}] \tag{16}$$

can be obtained from any linear combination of the two vector potentials

$$\mathbf{A_s} = b_0 \left[y \cos(\nu t) - x \sin(\nu t) \right] \mathbf{e_z},\tag{17a}$$

$$\mathbf{A_a} = b_0 z \left[\sin(\nu t) \mathbf{e_x} - \cos(\nu t) \mathbf{e_y} \right], \tag{17b}$$

and that each of this vector potential combination leads to a different particle dynamic as a result of a different inductive electric field $\mathbf{E} = -\partial \mathbf{A}/\partial t$. Furthermore, the orbit of a particle in a given combination of $\mathbf{A_s}$ and $\mathbf{A_a}$ can not be simply deduced from the particle orbits in $\mathbf{A_s}$ and $\mathbf{A_a}$ separately [90].

For each vector potential field, ion stability criteria depends on the two dimensionless parameters ν/Ω and b_0/B_0 , with $\Omega = |q|B_0/m$. Since $\Omega \propto m^{-1}$, any stability frontier which is not purely horizontal in the $(\nu/\Omega, b_0/B_0)$ plane offers opportunities for mass separation. For example, the stability diagram for the simple case $\mathbf{A_s}$, as derived in Ref. [90], is plotted in Fig. 8. For any rotating field amplitude $b_0 \neq 0$,

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59 60 separation based on mass can be done in two ways. Consider first case \mathcal{B} in Fig. 8. Here, as the ion mass decreases from m_h to m_l at constant driving frequency ν , ν/Ω decreases below the threshold for stability for a given b_0/B_0 . As a result the orbit of a heavy ion of mass m_h is radially stable, but, for the same parameters ν , b_0 and B_0 , a light ion of mass m_l is not. On the other hand, the orbit of a light ion of mass m_l is stable while the orbit of a heavy ion of mass m_h is not for case \mathcal{A} . Indeed, here ν/Ω decreases below the threshold for stability for a given b_0/B_0 as the ion mass increases from m_l to m_h . The possibility to choose which of heavy or light particles is radially confined could prove very useful for some applications, and addresses one of the limitations of the Archimedes filter [63] discussed earlier in this section.

In light of these results, rotating magnetic field configurations appear promising for mass separation applications. However, it should be stressed that the vector potential field **A** one will obtain for a given driving currents configuration might differ from the ideal case considered above. Take for example $\mathbf{A_s}$. Because the inductive electric field $-\partial \mathbf{A_s}/\partial t$ is along the static magnetic field lines, screening is expected to take place, and the effective $\mathbf{A_s}$ will then be a function of position [90]. Besides screening, collisional effects will further complexify the vector potential. However, the detailed impact of screening and collisional effects on the practical operating parameter space of these configurations, and in turn on the achievable throughput, remains to be studied.

4. Summary

Innovative separation technologies could offer ingenious solutions to important societal challenges. One example of innovative separation technology is plasma separation. Plasma separation stands out from conventional separation techniques by allowing separation at the elemental level based on physical properties. Once a plasma is made out of the mixture to be separated, all differential transport and confinement properties found in plasmas can be leveraged to produce separation. In essence, plasma separation is an extension of plasma confinement physics, but for which the focus is shifted from maximizing confinement to maximizing differential effects.

One particular physical criteria for separation in a plasma is atomic mass. Plasma mass separation could prove valuable for nuclear waste cleanup, nuclear spent fuel reprocessing and rare earth recycling. Although isotope separation motivated the development of a few plasma mass filtration concepts in the 1980s, most of these concepts feature limited throughput. Most

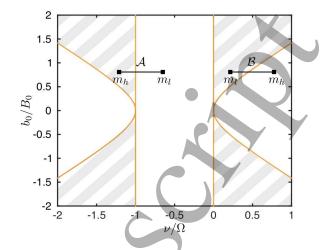


Figure 8. Stability diagram for the vector potential $\mathbf{A_s}$, from [90]. ν is the rotating field angular frequency, $\Omega = |q|B_0/m$ is the gyro-frequency and B_0 and b_0 are the axial and rotating magnetic field amplitudes, respectively. Hatched regions denote unstable regions, where ions are radially unconfined (the particle radial position diverges). m_l and m_h denote the mass of two ions $(m_l < m_h)$. Since $\Omega \propto m^{-1}$, there exists ν such that the light ion is confined while the heavy ion is unconfined (case \mathcal{B}), and reciprocally (case \mathcal{A}).

often, this limit results from constrains on plasma density set by collisions. Since new applications require processing large quantities of material, there is a need for developing new plasma mass separation concepts.

Although mass separation can be envisioned in many ways, rotating plasmas hold promise owing to centrifugal forces. Rotating plasmas can be used similarly to spinning gases or liquids to separate elements in plasma centrifuges. However, the uniqueness of plasmas lies in the fact that other forces can be leveraged in combination with centrifugal forces. This obviously includes electric and magnetic forces, but also mirror forces. A particle in a spinning gas column sees a mass dependent parabolic centrifugal potential. On the other hand, a charged particle in an $\mathbf{E} \times \mathbf{B}$ spinning plasma column (axial magnetic field, radial electric field) sees the same parabolic centrifugal potential plus an electric potential which depends on the applied potential radial profile. The extra control knob offered by the applied electric potential allows to confine radially light ions while deconfining heavy ions, or to separate light and heavy ions in distinct radial potential wells. Further means of control on particle dynamics can be obtained if abandoning the purely axial magnetic field topology. One solution then consists in combining magnetic pressure with variation of centrifugal potential along the magnetic field line to create mass differential confinement properties. Finally, another way to produce plasma rotation consists in using a rotating magnetic field. can in principle be achieved through different driving currents configurations, which each leads to different

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mass confinement capabilities.

The development of actual plasma mass filtering devices hinges on the ability to produce and control transverse electric fields for $\mathbf{E} \times \mathbf{B}$ rotating devices, or suitable potential vector fields for rotating magnetic field configurations, which have not yet been demonstrated on a practical device. A significant experimental effort is therefore needed to demonstrate these key elements, in particular for plasma parameters which are compatible with high-throughput separation applications. To the extent that perpendicular transport and rotation play a key role on performances in a large number of cross-field devices, it is anticipated that the results of this effort will benefit many applications besides mass separation.

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- [1] Sholl D S and Lively R P 2016 Nature 532 435-437 URL http://dx.doi.org/10.1038/532435a
- [2] National Research Council 1998 Separation Technologies for the Industries of the Future (National Academies Press) URL http://dx.doi.org/10.17226/6388
- [3] Oak Ridge National Laboratory 2005 Materials for separation technologies: Energy and emission reduction opportunities Tech. rep. URL http://www.dx.doi.org/ 10.2172/1218755
- [4] Eastland B and Gough W 1968 Bull. Am. Phys. Soc., Ser. II 1564
- [5] Dempster A J 1918 Phys. Rev. 11 316-325 URL http: //dx.doi.org/10.1103/physrev.11.316
- Lawrence E O 1958 Calutron system
- [7] Smith L P, Parkins W E and Forrester A T 1947 Phys. Rev. 72 989-1002 URL http://dx.doi.org/10.1103/ physrev.72.989
- [8] Parkins W E 2005 Phys. Today 58 45-51 URL http: //dx.doi.org/10.1063/1.199574
- [9] Alexeff I 1978 IEEE Transactions on Plasma Science 6 539-542 URL http://Dx.doi.org/10.1109/tps.1978. 4317160
- [10] Bonnevier B 1966 Ark. Fys. 33 255
- [11] Bonnevier B 1971 Plasma Phys. 13 763 URL https://doi. org/10.1088/0032-1028/13/9/007
- [12] Lehnert B 1971 Nucl. Fusion 11 485- URL http://dx.doi. org/10.1088/0029-5515/11/5/010
- [13] O'Neil T M 1981 Phys. Fluids 24 1447-1451 URL http: //dx.doi.org/10.1063/1.863565 [14] Lehnert B 1973 *Phys. Scr.* **7** 102 ISSN 1402-4896 URL
- https://doi.org/10.1088/0031-8949/7/3/002
- [15] Krishnan M, Geva M and Hirshfield J L 1981 Phys. Rev. Lett. 46 36-38 URL http://dx.doi.org/10.1103/ PhysRevLett.46.36
- [16] Del Bosco E, Simpson S W, Dallaqua R S and Montes A 1991 J. Phys. D: Appl. Phys. 24 2008- ISSN 0022-3727 URL http://dx.doi.org/10.1088/0022-3727/24/
- [17] James B W and Simpson S W 1976 Plasma Phys. 18 289 ISSN 0032-1028 URL https://doi.org/10.1088/ 0032-1028/18/4/004
- [18] Krishnan M 1983 Phys. Fluids 26 2676-2682 URL http: //dx.doi.org/10.1063/1.864460

- [19] Prasad R R and Krishnan M 1987 J. Appl. Phys. 61 4464-4470 URL http://dx.doi.org/10.1063/1.338407
- [20] Hirshfield J L, Levin L A and Danziger O 1989 IEEE Trans. Plasma Sci. 17 695-700 URL http://dx.doi.org/10. 1109/27.41184
- [21] Bittencourt J A and Ludwig G O 1987 Plasma Phys. Controlled Fusion 29 601 ISSN 0741-3335 URL https: //doi.org/10.1088/0741-3335/29/5/003
- [22] Fetterman A J and Fisch N J 2011 Phys. Plasmas 18 103503-8 URL http://dx.doi.org/10.1068/1.3646311
- [23] Grossman M W and Shepp T A 1991 IEEE Trans. Plasma Sci. 19 1114-1122 ISSN 0093-3813 URL http://dx.doi. org/10.1109/27.125034
- [24] Bokhan P A, Buchanov V V, Fateev N V, Kalugin M M and Kazaryan M A 2006 Laser Isotope Separation in Atomic Vapor (Wiley VCH Verlag GmbH) ISBN 3527406212
- [25] Dawson J M, Kim H C, Arnush D, Fried B D, Gould R W, Heflinger L O, Kennel C F, Romesser T E, Stenzel R L, Wong A Y and Wuerker R F 1976 Phys. Rev. Lett. 37 1547-1550 URL http://dx.doi.org/10.1103/ physrevlett.37.1547
- [26] Rax J M, Robiche J and Fisch N J 2007 Phys. Plasmas 14 043102-8 URL http://dx.doi.org/10.1063/1.2717882
- [27] Festeau-Barrioz M C and Sawley M L 1985 Plasma Phys. Controlled Fusion 27 641-651 URL http://dx.doi.org/ 10.1088/0741-3335/27/6/002
- [28] Eiedler-Ferrari N and Galvao R M O 1986 Plasma Phys. Controlled Fusion 28 515-525 URL http://dx.doi.org/ 10.1088/0741-3335/28/3/001
- [29] Hirshfield J 1976 Nucl. Instrum. Methods 139 119-123 URL http://dx.doi.org/10.1016/0029-554x(76) 90666-2
- [30] Chen F F 1991 Double helix: the dawson separation process From Fusion to Light Surfing, Lectures on Plasma Physics Honoring John M Dawson ed Katsouleas T (Addison-Wesley, New York)
- Feinberg R M and Hargrove R S 1993 Overview of uranium atomic vapor laser isotope separation Tech. Rep. UCRL-ID-114671 Lawrence Livermore National Laboratory
- [32] Jenson R J, Judd O P and Sullivan J A 1982 Separating isotopes with lasers Los Alamos Science No. 4
- [33] Noyes R 1996 Nuclear Waste Cleanup Technologies and Opportunities (William Andrew Pub) ISBN 081551381X
- [34] Friedman G H 2014 Special report management challenges at the department of energy? fiscal year 2015 Tech. Rep. DOE/IG-0924 U.S. Department of Energy, Office of Inspector General, Office of Audits and Inspections
- [35] Swanson J L 1993 Clean option: An alternative strategy for hanford tank waste remediation Tech. Rep. PNNL-8388 Pacific Northwest National Laboratory
- [36] Office of Science Workshop on Environmental Management 2016 Basic research needs for environmental management Tech. rep. U. S. Department of Energy URL http://science.energy.gov/~/media/bes/ pdf/reports/files/BRNEM_rpt.pdf
- [37] Toumanov Y N 2003 Plasma and High Frequency Processes for Obtaining and Processing Materials in the Nuclear Fuel Cycle (Fizmatlib)
- [38] Nash K L and Lumetta G J (eds) 2011 Advanced Separation Techniques for Nuclear Fuel Reprocessing and Radioactive Waste Treatment (Woodhead publishing)
- [39] ESNII Executive Board 2013 European sustainable nuclear industrial initiative - implementation plan 2013-15 Tech. rep. Sustainable Nuclear Energy Technology Platform
- [40] OECD Nuclear Energy Agency 2014 Technology roadmap update for generation iv nuclear energy systems Tech. rep. Generation IV International Forum
- [41] Magill J, Berthou V, Haas D, Galy J, Schenkel R, Wiese H W, Heusener G, Tommasi J and G Y 2003 Nuclear Energy 42 263-277

59 60

- [42] Blue Ribbon Commission on America's Nuclear Future 2012

 Reactor and Fuel Cycle Technology Subcommittee report

 to the full commission updated report (Washington,
 DC: Blue Ribbon Commission on America's Nuclear
 Future) URL http://forward.library.wisconsin.edu/
 catalog/ocn781423926
- [43] Mathur J N, Murali M S and Nash K L 2001 Solvent Extr. Ion Exch. 19 357–390 ISSN 0736-6299 URL http: //dx.doi.org/10.1081/SEI-100103276
- [44] Peterman D R, Greenhalgh M R, Tillotson R D, Klaehn J R, Harrup M K, Luther T A, Law J D and Daniels L M 2008 Separation of minor actinides from lanthanides by dithiophosphinic acid extractants In Proceedings of the 2008 Internation Solvent Extraction Conference (ISEC)
- [45] Morss L R, Edelstein N M and Fuger J (eds) 2011 The Chemistry of the Actinide and Transactinide Elements (Springer Netherlands) URL http://dx.doi.org/10. 1007/978-94-007-0211-0
- [46] Gelis A V and Lumetta G J 2014 Industrial & Engineering Chemistry Research 53 1624–1631
- [47] Barteková E and Kemp R 2016 Resources Policy 49 153– 164 ISSN 0301-4207 URL http://dx.doi.org/10.1016/ j.resourpol.2016.05.003
- [48] Charalampides G, Vatalis K, Karayannis V and Baklavaridis A 2016 IOP Conference Series: Materials Science and Engineering 161 012069 ISSN 1757-899X URL https://doi.org/10.1088/1757-899X/ 161/1/012069
- [49] Binnemans K, Jones P T, Blanpain B, Van Gerven T, Yang Y, Walton A and Buchert M 2013 J. Cleaner Prod. 51 1-22 ISSN 0959-6526 URL http://www.sciencedirect. com/science/article/pii/S0959652612006932
- [50] Gueroult R, Hobbs D T and Fisch N J 2015 J. Hazard. Mater. 297 153-159 ISSN 0304-3894 URL http://dx.doi.org/10.1016/j.jhazmat.2015.04.058
- [51] Gueroult R and Fisch N J 2014 Plasma Sources Sci. Technol. 23 035002- ISSN 0963-0252 URL http://dx. doi.org/10.1088/0963-0252/23/3/035002
- [52] Gueroult R, Rax J M and Fisch N J 2017 Submitted to Environmental Science and Technology
- [53] Bardez I 2004 Etude des caractéristiques structurales et des propriétés de verres riches en terres rares destinés au confinement des produits de fission et éléments à vie longue Ph.D. thesis Université Pierre et Marie Curie URL http://pastel.archives-ouvertes.fr/pastel-00001075
- [54] Grossbeck M L, Renier J P A and Bigelow T 2003 Development of improved burnable poisons for commercial nuclear power reactors Tech. Rep. NERI Project Number 99-0074 University of Tennessee final Report URL https://www.osti.gov/scitech/servlets/purl/820689
 [55] Timofeev A V 2014 Sov. Phys. Usp. 57 990- ISSN 1063-
- [55] Timofeev A V 2014 Sov. Phys. Usp. 57 990- ISSN 1063-7869 URL http://dx.doi.org/10.3367/UFNe.0184. 201410g.1101
- [56] Kaplan A F H 2014 Journal of Laser Applications 26 012007 ISSN 1042-346X URL http://dx.doi.org/10. 2351/1.4833936
- [57] Lieberman M A and Lichtenberg A J 1994 Principles of Plasma Discharge for Materials Processing (John Wiley & Sons)
- [58] National Research Council (ed) 2001 Research Needs for High-Level Waste Stored in Tanks and Bins at U.S. Department of Energy Sites: Environmental Management Science Program (The National Academies Press) URL http://www.nap.edu/catalog.php?record_id=10191
- [59] Zhil'tsov V A, Kulygin V M, Semashko N N, Skovoroda A A, Smirnov V P, Timofeev A V, Kudryavtsev E G, Rachkov V I and Orlov V V 2006 Atomic Energy 101 755-759 ISSN 1573-8205 URL http://dx.doi.org/10.

- 1007/s10512-006-0164-7
- [60] Davidson R C 2001 Physics of Nonneutral Plasmas (Imperial College Press)
- [61] Hestenes D 1998 New fondations for classical mechanics 2nd ed (Fundamental Theories of Physics vol 99) (Springer)
- [62] Ohkawa T and Miller R L 2002 Phys. Plasmas 9 5116-5120 URL http://dx.doi.org/10.1063/1.1523930
 [63] Freeman R, Agnew S, Anderegg F, Cluggish B, Gilleland J,
- [63] Freeman R, Agnew S, Anderegg F, Cluggish B, Gilleland J, Isler R, Litvak A, Miller R, O'Neill R, Ohkawa T, Pronko S, Putvinski S, Sevier L, Sibley A, Umstadter K, Wade T and Winslow D 2003 AIP Conf. Proc. 694 403-410 URL http://dx.doi.org/10.1063/1.1638067
- URL http://dx.doi.org/10.1063/1.1638067
 [64] Rax J M, Fruchtman A, Gueroult R and Fisch N J 2015
 Phys. Plasmas 22 092101 URL http://dx.doi.org/10.
 1063/1.4929791
- [65] Gueroult R, Rax J M and Fisch N J 2014 Phys. Plasmas 21 020701 URL http://dx.doi.org/10.1063/1.4864325
 [66] Kent G I, Jen N C and Chen F F 1969 Phys.
- [66] Kent G I, Jen N C and Chen F F 1969 Phys. Fluids 12 2140-2151 URL http://dx.doi.org/10.1063/ 1.1692323
- [67] Jassby D L 1972 Phys. Fluids 15 1590-1604 URL http: //aip.scitation.org/doi/abs/10.1063/1.1694135
- [68] Rosenbluth M N, Krall N A and Rostoker N 1962 Nucl. Fusion Suppl. Pt. 1 143
- [69] Chen F F 1966 Phys. Fluids 9 965-981 URL http://aip. scitation.org/doi/abs/10.1063/1.1761798
- [70] Gueroult R, Rax J M and Fisch N J 2017 Phys.

 Plasmas 24(8) 082102 URL http://dx.doi.org/10.
 1063/1.4994546
- [71] Ferraro V C A 1937 Mon. Not. R. Astron. Soc. 97 458-472
- [72] Volosov V 2006 Nucl. Fusion 46 820—ISSN 0029-5515 URL http://dx.doi.org/10.1088/0029-5515/46/8/007
- [73] Lehnert B 1974 Phys. Scr. **9** 189–192
- [74] Bekhtenev A A, Volosov V I, Pal'chikov V E, Pekker M S and Yudin Y N 1980 Nucl. Fusion 20 579–598
- [75] Fetterman A J and Fisch N J 2011 Phys. Plasmas 18 094503-3 URL http://dx.doi.org/10.1063/1.3631793
- [76] Gueroult R and Fisch N J 2012 Phys. Plasmas 19 122503-6 URL http://dx.doi.org/10.1063/1.4771674
- [77] Ochs I E, Gueroult R, Fisch N J and Zweben S J 2017 Phys. Plasmas 24 043503 URL http://dx.doi.org/10.1063/ 1.4978949
- [78] Lehnert B 1970 Phys. Scr. 2 106 ISSN 1402-4896 URL http://doi.org/10.1088/0031-8949/2/3/007
- [79] Tsushima A, Mieno T, Oertl M, Hatakeyama R and Sato N 1986 Phys. Rev. Lett. 56 1815–1818 URL http://dx. doi.org/10.1103/PhysRevLett.56.1815
- [80] Tsushima A and Sato N 1991 J. Phys. Soc. Jpn. 60 2665-2672 ISSN 0031-9015 URL http://dx.doi.org/ 10.1143/JPSJ.60.2665
- [81] Shinohara S and Horii S 2007 Jap. J. App. Phys. 46 4276- ISSN 1347-4065 URL http://dx.doi.org/10. 1143/JJAP.46.4276
- [82] Gueroult R, Evans E S, Zweben S J, Fisch N J and Levinton F 2016 Plasma Sources Sci. Technol. 25 035024 ISSN 0963-0252 URL http://dx.doi.org/10. 1088/0963-0252/25/3/035024
- [83] Fetterman A J and Fisch N J 2009 Plasma Sources Sci. Technol. 18 045003 ISSN 0963-0252 URL http://dx. doi.org/10.1088/0963-0252/18/4/045003
- [84] Fisch N J and Rax J M 1992 Phys. Rev. Lett. 69 612-615 URL https://doi.org/10.1103/PhysRevLett.69.612
- [85] Fetterman A J and Fisch N J 2008 Phys. Rev. Lett. 101 205003- URL http://dx.doi.org/10.1103/ PhysRevLett.101.205003
- [86] Stepanov V G, Zakharcthenko V F and Bezel' V S 1958 JETP 34 512 URL http://www.jetp.ac.ru/cgi-bin/dn/e_007_02_0353.pdf
- [87] Soldatenkov T R 1966 Sov. Phys. Tech. Phys. 11 179

Mass differential effects in magnetized rotating plasmas for separation applications

- [88] Fisch N J and Watanabe T 1982 Nuclear Fusion ${\bf 22}$ 423
- [89] Hugrass N W and Jones I R 1983 J. Plasma Physics 29 155
- [90] Rax J M and Gueroult R 2016 J. Plasma Phys. 82(5) 595820504 URL http://dx.doi.org/10.1017/ S0022377816000878